L Number	Hits	Search Text		Time stamp
- Number	2	20030004103.did.	USPAT;	2003/11/03
	İ	i İ	US-PGPUB; EPO; JPO;	02:36
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-	2	20020028772.did.	USPAT;	2003/10/23
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	1	1343876.did. and metabolism	DERWENT USPAT;	2002/10/22
!	ļ	1343070.dld. and metapolism	US-PGPUB;	2003/10/23 06:37
İ	; !	!	EPO; JPO;	
-	2	9853050.did. and peptide	USPAT;	2003/10/23
			US-PGPUB; EPO; JPO;	06:38
:			DERWENT	0000/11/45
-	8	"HJ loop" and modulate and "serine/threonine kinase"	'USPAT; US-PGPUB;	2003/11/03 02:36
	į		EPO; JPO;	!
_	7	"HJ loop" and modulating and	USPAT;	2003/11/03
		"serine/threonine kinase" and method	US-PGPUB; EPO; JPO;	02:37
			DERWENT	
· -	7	"HJ loop" and modulate and "serine/threonine kinase" and method	USPAT; US-PGPUB;	2003/11/03 02:37
			EPO; JPO;	
_	17	""HJ loop" and modulate	DERWENT USPAT;	2003/11/03
	·		US-PGPUB; EPO; JPO;	02:38
			DERWENT	
-	18	"HJ loop"	USPAT; US-PGPUB;	2003/11/03 02:46
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-	1485	 	DERWENT USPAT;	2003/11/03
			US-PGPUB; EPO; JPO;	02:41
			DERWENT	
<u> </u>	454	"serine/threonine kinase" and loop	USPAT; US-PGPUB;	2003/11/03 02:41
	İ	I	EPO; JPO;	
-	9	"serine/threonine kinase" and "HJ" and	DERWENT USPAT;	2003/11/03
		loop	US-PGPUB; EPO; JPO;	02:42
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:-	11598	modulation and kinase	USPAT; US-PGPUB;	2003/11/03
	i		EPO; JPO;	
-	1537	modulation SAME kinase	DERWENT USPAT;	2003/11/03
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; 	!		DERWENT	
-	19	modulation SAME kinase SAME loop	USPAT; US-PGPUB;	2003/11/03 02:52
:			EPO; JPO;	
_	0	"Val-Met-Thr-Gly-Glu-Leu-Pro-Phe"	DERWENT USPAT;	2003/11/03
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		<u> </u>	DERWENT	<u></u>

		Val and Met and Thr and Gly and Glu and Leu and Pro and Phe	USPAT; US-PGPUB; EPO; JPO; DERWENT	2003/11/03
-	19600	Val SAME Met SAME Thr SAME Gly SAME Glu SAME Leu SAME Pro SAME Phe	USPAT; US-PGPUB; EPO; JPO;	2003/11/03 02:54
: -	0	"V-M-T-G-Q-L-P-F"	DERWENT USPAT; US-PGPUB; EPO; JPO;	2003/11/03
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Dialog level 03.04.00D
Last logoff: 03nov03 04:02:32
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          *** ANNOUNCEMENT ***
--File 654 - US published applications from March 15, 2001 to the
present are now online. Please see HELP NEWS 654 for details.
--File 581 - The 2003 annual reload of Population Demographics is
complete. Please see Help News581 for details.
--File 990 - NewsRoom now contains February 2003 to current records.
File 992 - NewsRoom 2003 archive has been newly created and contains
records from January 2003. The oldest months's records roll out of
File 990 and into File 992 on the first weekend of each month.
To search all 2003 records BEGIN 990, 992, or B NEWS2003, a new
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--Connect Time joins DialUnits as pricing options on Dialog.
See HELP CONNECT for information.
                   *** --SourceOne patents are now delivered to your email inbox
as PDF replacing TIFF delivery. See HELP SOURCE1 for more
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-- Important news for public and academic
libraries. See HELP LIBRARY for more information.
-- Important Notice to Freelance Authors--
See HELP FREELANCE for more information
NEW FILES RELEASED
***Emergency Room (File 454), Hospital Inpatient Profiles (File 462),
   and Hospital Outpatient Profiles (File 463)
***World News Connection (File 985)
***Dialog NewsRoom - 2003 Archive (File 992)
***TRADEMARKSCAN-Czech Republic (File 680)
***TRADEMARKSCAN-Hungary (File 681)
***TRADEMARKSCAN-Poland (File 682)
UPDATING RESUMED
                   * * *
RELOADED
***Population Demographics - (File 581)
***CLAIMS Citation (Files 220-222)
                     *** DIALOG HOMEBASE(SM) Main Menu ***
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  1. Announcements (new files, reloads, etc.)
  2. Database, Rates, & Command Descriptions
  3. Help in Choosing Databases for Your Topic
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  5. Product Descriptions
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  6. DIALOG(R) Document Delivery
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/H = Help
                          /L = Logoff
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 service. Enter a BEGIN command plus a file number to search a database
(e.g., Bl for ERIC).
B 155, 73, 344, 358, 65, 35
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 *File 155: On 13 November, Medline will temporarily stop updating with
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      S2
S S1 AND "HJ"
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DIALOG(R) File 155:MEDLINE(R)
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15524478
                      PMID: 12763936
           22840662
 Induction of pro-angiogenic signaling by a synthetic peptide derived from
 the second intracellular loop of S1P3 (EDG3).
         Tamar;
                   Tsirulnikov
                                Lilia; Reuveni Hadas; Yarnitzky Talia;
Ben-Sasson Shmuel A
  Keryx Biopharmaceuticals, Jerusalem, Israel.
  Blood (United States)
                        05 22 2003, 102 (6) p2099-107, ISSN 0006-4971
Journal Code: 7603509
  Document type: Journal Article
  Languages: ENGLISH
  Main Citation Owner: NLM
  Record type: Completed
  The G-protein-coupled receptors of the endothelial differentiation gene
(EDG) family mediate pro-angiogenic activities, such as endothelial cell
proliferation, chemotaxis, and vessel morphogenesis. We synthesized and
tested the effects of a 9-amino acid peptide (KRX-725), derived from the
second intracellular loop of S1P3 (EDG3). KRX-725 mimics the effects of
sphingosine 1-phosphate (S1P), the natural ligand of S1P3, by triggering a
Gi-dependent MEK-ERK (mitogen-activated protein
                                                        kinase kinase and
extracellular signal-regulated kinase) signal transduction pathway. Using
aortic rings as an ex vivo model of angiogenesis, vascular sprouting was
assessed in the presence of KRX-725 or S1P. KRX-725 induced extensive and
dense vascular sprouts, which contain an elaborated organization of endothelial and smooth muscle layers, including lumen formation. When
KRX-725 or S1P was combined with proangiogenic factors, such as basic
fibroblast growth factor (bFGF), stem cell factor, or vascular endothelial
growth factor, the effect was synergistic, leading to further enhancement
of vascular sprouting. KRX-725 also initiated neovascularization in a mouse
corneal pocket assay in vivo and showed synergism with bFGF. The specificity of KRX-725 was demonstrated via peptide-induced receptor
internalization of S1P3 but not S1P1. The ability of a short peptide to
stimulate extensive angiogenesis and to synergize with pro-angiogenic
factors suggests that KRX-725 may serve as a useful agent in treating
pathologic conditions
                         such as peripheral vascular disease, cardiac
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7/3,AB/2 (Item 1 from file: 73)

DIALOG(R) File 73: EMBASE

ischemia, or tissue grafts.

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12265714 EMBASE No: 2003375236

Induction of pro-angiogenic signaling by a synthetic peptide derived from the second intracellular loop of S1PSUB3 (EDG3)

Licht T.; Tsirulnikov L.; Reuveni H.; Yarnitzky T.; Ben-Sasson S.A. S.A. Ben-Sasson, Dept. of Exp. Med./Cancer Research, Hebrew

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Univ.-Hadassah Medical School, POB 12272, Jerusalem Israel AUTHOR EMAIL: muli@md.huji.ac.il Blood ( BLOOD ) (United States) 15 SEP 2003, 102/6 (2099-2107) CODEN: BLOOA ISSN: 0006-4971 DOCUMENT TYPE: Journal; Article LANGUAGE: ENGLISH SUMMARY LANGUAGE: ENGLISH NUMBER OF REFERENCES: 54
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The G-protein-coupled receptors of the endothelial differentiation gene (EDG) family mediate pro-angiogenic activities, such as endothelial cell proliferation, chemotaxis, and vessel morphogenesis. We synthesized and tested the effects of a 9-amino acid peptide (KRX-725), derived from the second intracellular loop of S1PSUB3 (EDG3). KRX-725 mimics the effects of sphingosine 1-phosphate (S1P), the natural ligand of S1P3, by triggering a GSUBi-dependent MEK-ERK (mitogen-activated protein kinase kinase and extracellular signal-regulated kinase) signal transduction pathway. Using aortic rings as an ex-vivo model of angiogenesis, vascular sprouting was assessed in the presence of KRX-725 or S1P. KRX-725 induced extensive and dense vascular sprouts, which contain an elaborated organization of endothelial and smooth muscle layers, including lumen formation. When KRX-725 or S1P was combined with pro-angiogenic factors, such as basic fibroblast growth factor (bFGF), stem cell factor, or vascular endothelial growth factor, the effect was synergistic, leading to further enhancement of vascular sprouting. KRX-725 also initiated neovascularization in a mouse corneal pocket assay in vivo and showed synergism with bFGF. The specificity of KRX-725 was demonstrated via peptide-induced receptor internalization of S1 P3 but not S1PSUB1. The ability of a short peptide to stimulate extensive angiogenesis and to synergize with pro-angiogenic factors suggests that KRX-725 may serve as a useful agent in treating pathologic conditions such as peripheral vascular disease, cardiac ischemia, or tissue grafts. (c) 2003 by The American Society of Hematology. COST

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